

REMARKS

Entry of the amendments is respectfully requested. Claims 17, 7, and 24 are currently amended. New claim 55 has been added. Claims 1, 7-25, and 28-55 are pending in the application. Favorable reconsideration and allowance of this application is respectfully requested in light of the foregoing amendments and the remarks that follow.

1. Examiner Interview

The applicant thanks the Examiner for the helpful interview held on October 10, 2006. In the interview, the functional language and the percent identity in the claims were discussed. For the proposed functional language, the Examiner made two proposals. First, he suggested “wherein the polypeptide is a marker of pluripotent cells.” Second, the Examiner proposed “wherein the polypeptide can serve as a marker of germ cell competence in embryonic germ cell precursors.” The Examiner indicated that the first proposal would require an RCE because it was broader, whereas he and his primary approved of the second proposal, but would require the SPE's approval. Applicant has amended the claims based on the first proposal and files an RCE herewith.

With respect to the percent identity, the Examiner suggested that claim 1 should be amended to require 95% identity. This suggestion was based on Example 14 of the “Revised Interim Written Description Guidelines Training Materials.” The Examiner’s proposals are discussed in more detail below.

2. Rejection Under § 112, First Paragraph

Claims 1, 7, and 24 were rejected under 35 U.S.C. § 112, ¶ 1, as failing to comply with the written description requirement. The applicant believes that this requirement was satisfied prior to the current amendments. Nevertheless, the claims have been amended, but only for the purpose of expediting prosecution. The amendments in no way represent a concession that the invention is fairly limited in this way.

As noted above, applicant has chosen amendment language based on the Examiner’s broader proposal. Support for the amendments, i.e., “the polypeptide can serve as a marker of pluripotent cells,” and the fact that the claims satisfy the written description requirement is found

at paragraphs 71 and 309 of the published patent application. Additional support can be found in the attached publications, Geijsen N., et al., *Derivation of Embryonic Germ Cells and Male Gametes from Embryonic Stem Cells*, Nature 427:148-153 (attached as Exhibit A) and Gordeeva, O., et al., *Differentiation of Embryonic Stem Cells after Transplantation into Peritoneal Cavity of Irradiated Mice and Expression of Specific Germ Cell Genes in Pluripotent Cells*, Trans. Proc. 2005, 37(1): 295-298 (attached as Exhibit B). Both of these references confirm that expression of GCR1/Fragilis is not limited to pluripotent germ cell precursors. Instead, GCR1/Fragilis is also expressed in other pluripotent cells, such as mouse ES cells. For example, Gordeeva et al. states that "Expression of pluripotent and germ-cell specific genes oct4, nanog, stella and fragilis was evaluated. . . . [A]ll four genes were expressed in the pluripotent cells. (see page 297, second column, last paragraph). Likewise, Figure 1 of Geijsen et al. clearly shows that Fragilis is expressed in mouse ES cells (see left hand lane of Figure 1). Thus, given that in the art, (1) the model pluripotent cell is considered to be the ES cell and (2) fragilis is acknowledged as a marker for ES cells as well as other pluripotent cells, such as embryonic germ cell precursors, there is ample support for the amendments. Hence, there is no reason to require the Examiner's narrower, second proposal as it would fly in the face of the currently understood scientific wisdom, as demonstrated in both Gordeeva et al. and Geijsen et al.

For the percent identity of claim 1, the Examiner suggests narrowing to it 95%. This was based on Example 14 of the "Revised Interim Written Description Guidelines Training Materials," which has a claim limitation of 95% identity. However, this is not a general rule. Instead, it is clearly specific to and based on the specification of that example:

Analysis: A review of the full content of the specification indicates that a protein having SEQ ID NO: 3 or variants having 95% identity to SEQ ID NO: 3 and having catalytic activity are essential to the operation of the claimed invention.

In the instant application, the specification supports a 90% identify. See, e.g., paragraphs 9, 62, and 221. Furthermore, the 90% identity defines over the prior art. As such, one skilled in the art would conclude that based on the 90% identity and the functional limitations of the claims, the applicant was in possession of the necessary common attributes possessed by the members of the genus of polypeptide encompassed by the claim. Thus, the written description requirement is satisfied.

In light of the amendments to the claims and the foregoing argument, withdrawal of the rejection of claims 1, 7, and 24 under 35 U.S.C. § 112, ¶ 1 is requested.

3. Withdrawal of the Rejection Under § 103

The withdrawal of the rejection of claim 1 under 35 U.S.C. § 103(a) is noted with thanks.

4. New Claim

New claim 55 has been added and recites "An isolated GCRI polypeptide comprising the sequence set forth in SEQ ID NO: 2." This claim is believed to be in condition for allowance for at least the reason that the other claims are believed to be in condition for allowance.


CONCLUSION

It is submitted that the pending claims are in compliance with 35 U.S.C. § 112 and each define patentable subject matter. A Notice of Allowance is therefore respectfully requested.

The Director is authorized to charge Deposit Account No. 23-2053 for the fee associated with a three -month extension, which the Applicants hereby request, and the fee associated with one additional independent claim. No other fee is believed to payable with this response. The Director is authorized to charge Deposit Account No. 23-2053 for any other fee associated with this response. The Examiner is invited to contact the undersigned by telephone if it would help expedite matters.

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expand (Supplementary Information). It is important to recognize that further work is required to establish empirically how the absolute and proportional area losses of individual species (in other words, the type of data from climate envelope projections) are related to extinction risk. As yet, no agreed standard method exists for such calculations: assumptions and uncertainties inherent in the three methods will be considered in detail elsewhere.

Extinction probability estimates were not available for all scenarios in every region/taxon, so means of scenarios were calculated after using a least-squares analysis of variance model to impute missing values. Region/taxon mean probabilities of extinction for each scenario were logit-transformed and a three-way analysis of variance was fitted (region/taxon \times climate scenario \times dispersal scenario; weighted by $\sqrt{N_{\text{species}}}$ per region/taxon study). The fitted model was used to impute expected values of the probability of extinction for those region/taxon and scenario combinations for which direct estimates were not available. Scenario means were then calculated from the combined direct estimates and imputed values, using $\sqrt{N_{\text{species}}}$ for each region/taxon as weights.

Red Data Book criteria

Each species is assigned to a threat category¹⁶, or classified 'Not Threatened' (0% risk), depending on the projected decline in area over 50 or 100 years (Supplementary Information) and the final distribution area. Existing areas were considered, so we present only the extra extinction attributable to climate change. Logit-transformed three-way analysis of variance was used to estimate extinction risks for empty cells, as with the species-area approaches.

Extinct: species with a projected future area of zero (100% of species assumed to be committed to eventual extinction).

Critically endangered: projected future distribution area $< 10 \text{ km}^2$, or decline by $> 80\%$ in 50 years (species assigned a 75% chance of extinction¹⁶).

Endangered: projected area $10\text{--}500 \text{ km}^2$, or $50\text{--}80\%$ decline in 50 years (species assigned a 35% chance of extinction¹⁶).

Vulnerable: projected area $500\text{--}2,000 \text{ km}^2$, or $> 50\%$ decline in 100 years on the basis of linear extrapolation of 50-year projection (species assigned a 15% chance of extinction¹⁶).

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Authors' contributions The fourth and subsequent authors are alphabetically arranged and contributed equally.

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Derivation of embryonic germ cells and male gametes from embryonic stem cells

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Egg and sperm cells (gametes) of the mouse are derived from a founder population of primordial germ cells that are set aside early in embryogenesis. Primordial germ cells arise from the proximal epiblast, a region of the early mouse embryo that also contributes to the first blood lineages of the embryonic yolk sac¹. Embryonic stem cells differentiate *in vitro* into cystic structures called embryoid bodies consisting of tissue lineages typical of the early mouse embryo^{2,3}. Because embryoid bodies sustain blood development, we reasoned that they might also support primordial germ cell formation. Here we isolate primordial germ cells from embryoid bodies, and derive continuously growing lines of embryonic germ cells. Embryonic germ cells show erasure of the

methylation markers (imprints) of the *Igf2r* and *H19* genes, a property characteristic of the germ lineage. We show that embryoid bodies support maturation of the primordial germ cells into haploid male gametes, which when injected into oocytes restore the somatic diploid chromosome complement and develop into blastocysts. Our ability to derive germ cells from embryonic stem cells provides an accessible *in vitro* model system for studies of germline epigenetic modification and mammalian gametogenesis.

We differentiated embryonic stem (ES) cells according to our standard methods⁴, and isolated messenger RNA from whole embryoid bodies (EBs) at several time points. We used polymerase chain reaction with reverse transcription (RT-PCR) to detect expression of genes implicated in ES cell pluripotency (the POU domain transcription factor *Oct4*) and germ cell development, including *stella* and *fragilis* (*Fgls*)⁵, and a set of genes that are exclusively expressed in the germ line and are absent from somatic tissues (*Dazl*, *Piwil2*, *Rnf17*, *Rnh2*, *Tdrd1* and *Tex14*)^{6–9}. All of these genes were expressed in undifferentiated ES cells, and a subset underwent rapid extinction with EB formation (Fig. 1). *Rnh2*, *Tdrd1* and *Tex14* decreased to undetectable levels very early in EB development (day 3–4), suggesting efficient differentiation and commitment to distinct cell fates. *Stella* and *Fgls* expression declined immediately on EB formation, but low levels persisted over the course of EB differentiation.

Expression of the surface antigen SSEA1, a marker of pluripotent ES cells, also wanes on EB development, but rare SSEA1 positive

(SSEA1⁺) cells persist in differentiated EBs¹⁰. We differentiated ES cells carrying an *Oct4*-promoter-driven green fluorescent protein (GFP) reporter gene¹¹, and observed a similar overall decrease in GFP⁺ cell populations upon EB differentiation, as well as a persistent rare population of GFP⁺ cells (data not shown). Although these might represent residual undifferentiated ES cells, SSEA1 and *Oct4* expression are also features of primordial germ cells (PGCs). We thus sought to determine whether these SSEA1⁺/Oct4⁺ cells within EBs represented residual undifferentiated ES cells or true PGCs.

There is a lack of markers that can suitably distinguish between ES cells and PGCs; however, retinoic acid acts to rapidly differentiate ES cells while stimulating proliferation of PGCs, and can therefore be used to distinguish between these two cell populations¹². We plated ES cells or cells derived from EBs onto a mouse embryonic feeder (MEF) cell layer and cultured the cells for 7 days in the presence of 2 μ M retinoic acid (Fig. 2a). We then quantified the residual SSEA1⁺ cells. In retinoic acid-treated ES cells, greater than 99% of cells extinguished SSEA1 expression. In contrast, in retinoic acid-treated cultures of EB-derived cells, a significant percentage of SSEA1⁺ cells persisted and expanded modestly in culture, with an apparent wave of formation peaking at around day 5 (Fig. 2b). Cells that differentiated for 5 days in retinoic acid were fixed and stained for alkaline phosphatase. Very few cells positive for alkaline phosphatase persisted in the retinoic acid-treated ES cell cultures (Fig. 2c, left panel). In contrast, large colonies of cells positive for alkaline phosphatase surrounded by motile cells that resembled migratory PGCs were abundant in retinoic acid-treated cultures of EB-derived cells (Fig. 2c, right panel). This differential effect of retinoic acid strongly suggested that the SSEA1⁺ population of cells from EBs were PGCs.

To obtain conclusive evidence that the retinoic acid-resistant ES-like colonies were indeed PGCs, we analysed whether these cells manifested erasure of epigenetic imprints; this is a unique property of PGCs that is maintained in PGC-derived embryonic germ cells¹³. Imprinting of the *Igf2r* gene is determined by parental origin, with expression only from the maternal allele¹⁴. A specific region of the *Igf2r* gene has been identified, differentially methylated region 2 (DMR2), which is hypermethylated only on the maternally inherited allele¹⁵. We isolated SSEA1⁺ cells from EBs at different time points and cultured the cells for 7 days in the presence of retinoic acid. Individual retinoic acid-resistant colonies were isolated and expanded on gelatinized tissue culture plastic in the presence of leukaemia inhibitory factor (LIF), stem cell factor (SCF) and basic fibroblast growth factor; these are conditions that support the derivation of embryonic germ cells^{16,17}. We analysed the methylation status of DMR2 in independent clones of the parental ES cell line and in candidate embryonic germ clones by restriction digestion of the genomic DNA with *PvuII* and the methylation-sensitive enzyme *MluI*. Independent ES cell clones demonstrated a somatic methylation profile in which only one allele was digested (Fig. 2d, top panel, lanes 1–5). Most of the day 4 EB-derived embryonic germ cell clones displayed a similar somatic imprinting profile (Fig. 2d, top panel, lanes 6–9), with the exception of one clone in which methylation was erased (Fig. 2d, top panel, lane 10), as demonstrated by digestion of both alleles. Notably, at day 7 of EB development, 6 of 7 embryonic germ-like cells showed an unmethylated pattern (Fig. 2d, top panel, lanes 11–16), and at day 10 of EB development all embryonic germ clones had lost imprinting of the *Igf2r* gene. Similar results were obtained showing erasure of the methylation marker at the *H19/Igf2* locus (Fig. 2d, bottom panel). These data demonstrate that the EB-derived PGCs display phenotypic and biological properties of PGCs developing *in vivo*.

We then used immunomagnetic bead sorting to isolate SSEA1⁺ candidate PGCs from differentiated EBs, and used RT-PCR to analyse expression of germ-cell-specific markers. *Oct4* was

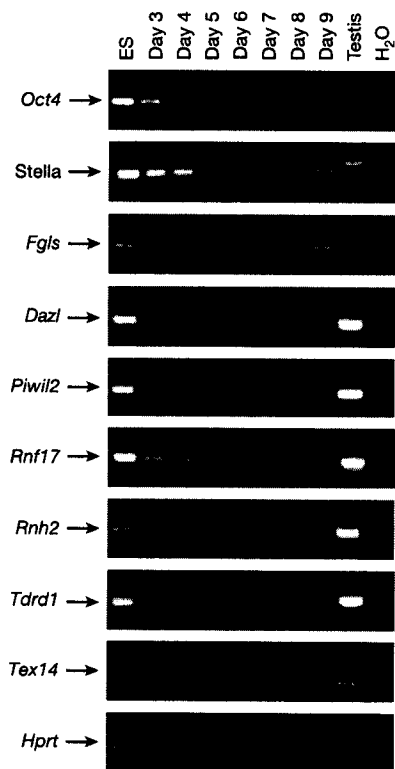


Figure 1 Expression of germ-cell-specific genes during EB development. RNA was prepared from ES cells, day 3–9 EBs and adult testis, and processed for RT-PCR. *Hprt* served as a control. *Stella* is expressed at high levels in primordial germ cells and oocytes but is almost absent from adult testis⁵. *Fgls* is expressed in the cells surrounding the germ cells⁵. *Dazl* is expressed only in germ cells in both testis and ovary^{7,29}. *Piwil2*, *Rnf17*, *Rnh2*, *Tdrd1* and *Tex14* are germ-cell-specific and expressed in the male gonad⁶.

expressed in the SSEA1⁺ population throughout EB development (Fig. 3a, left panel). *Oct4* expression was almost undetectable in the SSEA1-negative (SSEA1⁻) fraction, demonstrating the effectiveness of the SSEA1 selection (Fig. 3a, right panel). *Tex14* and *Rnh2*, which demonstrated a rapid downregulation on EB differentiation in the whole EB population, became undetectable in the purified SSEA1⁺ fraction of early EBs (days 3, 4; Fig. 3b). At day 5 expression levels of these genes rose, and by day 6 they reached a level comparable to ES cells. Although less marked than *Tex14* and *Rnh2*, expression of *Piwi2* and *Dazl* follows a similar pattern of increased expression over time in SSEA1⁺ differentiated EB-derived cell populations. This suggests that the developing EB supports a cellular environment similar to the early embryonic microenvironment in which PGCs are found.

To determine whether the PGCs arise in a defined region of the EB, we used immunohistochemistry to simultaneously visualize CD41⁺ haematopoietic cells¹⁸ and SSEA1⁺ germ cells in cryosections of 7-day-old EBs (Fig. 3c). Similar to the developing embryo, the SSEA1⁺ PGCs in the developing EB exist in close juxtaposition to the cells of the developing haematopoietic system. The colocalization of nascent blood and germ cell populations in the peripheral zones of the developing EB raises interesting questions about their clonal origins and the microenvironment that specifies their distinct cell fates.

We next investigated whether the EB-derived PGCs undergo further differentiation into functional gametes. We analysed the expression of *Sry*, a male gene that determines germ cell fate. *Sry*

expression was first detected in day 5 EBs, heralding initiation of a male germ cell developmental programme (Fig. 4a). Germ cell nuclear factor (*Gcnf*) has a role in confining *Oct4* expression to the germ line¹⁹. We observed transient expression of *Gcnf* starting at day 7 and peaking at around day 11, suggesting a temporal window during which the germ cells become fully specified. At about day 11 of EB differentiation we found a strong upregulation of acrosin and haprin, genes tightly associated with male germ cell development (Fig. 4b). Acrosin is part of the acrosomal complex transcribed as proacrosin in the diploid germ cell population²⁰, whereas haprin is a member of the RING finger-B box-coiled-coil family of transcription factors, which have a role in spermatogenesis and the formation of germ cell tumours^{21–23}. Our observation of upregulation of genes associated with male germ cell maturation prompted us to investigate the presence of stromal supporter cells. Indeed, we detected the message for the luteinizing hormone/gonadotropin receptor (LH-R) as well as müllerian inhibiting substance (MIS); these are markers of Leydig and Sertoli cells, respectively. We failed to detect expression of zona pellucida proteins Zp1 and Zp2, whereas Zp3 is expressed in ES cells and early EBs (Fig. 4b), suggesting that within the context of EB differentiation, the default programme of female gametogenesis is suppressed. Indeed when comparing the expression of two genes specific for male germ cell differentiation, *AZI* and ret finger protein (*Rfp*), we observed exclusive expression of these genes in EBs derived from male (XY) but not female (XX) ES cells (Fig. 4c).

To investigate whether male germ cells undergo meiosis in the

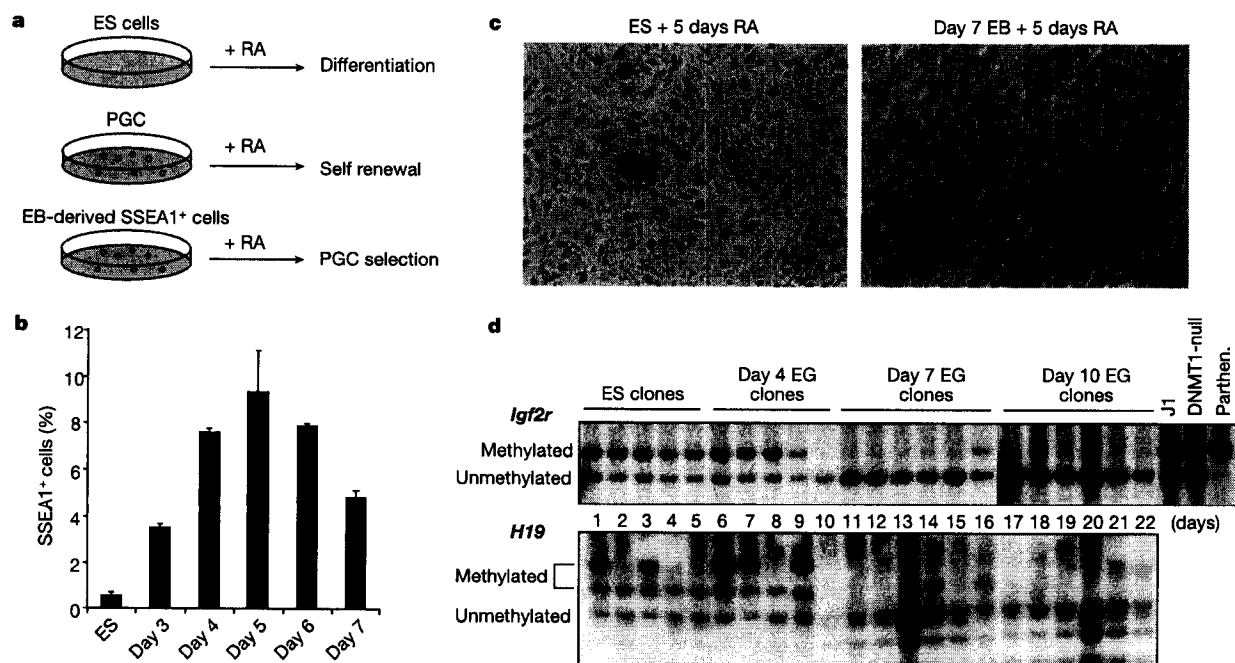


Figure 2 Development of primordial germ cells in the differentiating EB. **a**, Effects of treatment with retinoic acid (RA) on EB-derived cells. Retinoic acid differentiates ES cells (top panel) but supports PGCs¹² (middle panel). Culture of EB-derived SSEA1⁺ cells in the presence of retinoic acid selects for PGCs that retain germ cell marker expression (bottom panel). **b**, SSEA1⁺ cells were isolated from EBs of different ages by immunomagnetic beads, cultured in retinoic acid for 7 days, and SSEA1 expression was analysed by flow cytometry. The percentage of GFP⁺ ES and EB-derived cells expressing SSEA1 is plotted ($n = 3$). **c**, Alkaline phosphatase (AP) expression on ES cells (left panel) and day 7 EB-derived cells (right panel) after 5 days of culture in retinoic acid. The day 7 EB-derived cells formed large colonies of cells positive for alkaline phosphatase that resembled PGCs

on gross inspection¹⁷. Cells positive for alkaline phosphatase migrated out of the large colony (arrowheads). **d**, Imprint erasure of *Igf2r* and *H19*/*Igf2* loci. Top panel, DMR2 region of *Igf2r* locus. Southern analysis of genomic DNA from individual ES and EB-derived embryonic germ clones digested with *PvuII* and *MluI* is shown. Bottom panel, *H19*/*Igf2* locus. The same clones as in the top panel were digested with *SacI* and *HhaI*. The day of EB development at which individual embryonic germ clones were derived is indicated. Control lanes include J1 ES cells (somatic methylation profile), methylation-deficient DNMT1^{null} ES cells³⁰ and parthenogenetic (parthen.) ES cells (showing a maternal methylation pattern on both *Igf2* alleles).

context of the EB, we immunostained cell populations with an antibody that specifically recognizes male meiotic germ cells (FE-J1²⁴), and analysed these cells for DNA content using the fluorescent DNA-binding dye Hoechst 33342. In the testes of adult mice, both FE-J1⁺ and haploid (1C) cells are readily identified (Fig. 4d, left column of top and middle panels). Analysing the FE-J1⁺ cell population for DNA content shows that this marker of male germ cells recognizes a minor diploid (2C) population of primary spermatocytes and a predominant population of cells with a haploid (1C) DNA complement, as reported²⁴ (Fig. 4d, bottom panel, left). In the EB cell suspension, haploid (1C) cells are discernable but obscured by a range of apoptotic cells with sub-2C DNA content (Fig. 4d, middle panel, right). However, when only the cells positive for the FE-J1 antibody are analysed, a distinct haploid (1C) population is observed, representing the predominant class of FE-J1-staining cells (Fig. 4d, bottom panel, right). The low proportion of antibody-positive cells (0.01%) and the relatively high ratio between diploid (2C) primary spermatocytes and haploid (1C) cells suggests that meiosis is highly inefficient in EBs. This experiment, however, demonstrates that the EB microenvironment

is permissive for male germ cell development and meiotic maturation.

We further analysed the FE-J1⁺ cells by immunofluorescence microscopy. The FE-J1 antibody recognizes the anterior acrosome on early and late pachytene spermatocytes and on round spermatids²⁴. As can be seen in Fig. 4e, FE-J1⁺ cells isolated from testes (top panel) or from day 20 EBs (bottom panel) demonstrate polarized apical perinuclear staining, indicating that the EB-derived cells have a similar morphology to testis-derived haploid cells, and possibly represent round spermatids.

Finally, we investigated the biological function of the EB-derived haploid cells and assayed their capacity to fertilize oocytes. We isolated FE-J1⁺/GFP⁺ haploid cells from day 20 EBs by flow cytometry, and performed intracytoplasmic injection into recipient oocytes. Five separate microinjection experiments performed in two independent laboratories produced comparable results. Approximately 50% of the injected oocytes supported cleavage to the 2-cell stage ($n = 125$), with 20% displaying progression to blastocysts. Figure 4f shows four representative blastocysts expressing the GFP transgene, indicating successful complementation of the oocyte genome by the EB-derived haploid cells. Furthermore, fluorescence *in situ* hybridization (FISH) using probes directed against an autosomal locus and markers on the X and Y chromosome demonstrate a normal diploid chromosome complement and expected ratios of male (XY) and female (XX) embryos (Fig. 4g). Efforts are underway to determine whether embryos arising from fertilization with EB-derived male gametes will develop normally after uterine transfer.

Germ cell development remains a largely unexplored but fascinating process of cell fate specification. Germ cells represent a privileged class of cells, given responsibility for perpetuating pluripotency and ensuring propagation of the gene pool. The genetic mechanisms that account for maintenance of pluripotency and restrict somatic differentiation are beginning to yield to molecular analyses⁵, but an *in vitro* model system that recapitulates germ cell specification will greatly facilitate these studies.

Recently, Schöler and colleagues reported the generation of oocytes from mouse ES cells in culture²⁵. The reported differentiation happened spontaneously over a period of nearly 50 days. In contrast to our method of ES cell differentiation into EBs, Schöler and colleagues used bulk two-dimensional differentiation on tissue culture plastic, in which both male and female lines of ES cells yielded oocytes. Given our observation of male germ cell development, we speculate that EBs may preserve more of the tissue organization reflective of the embryonic gonadal ridge, thereby enabling male germ lineage specification. In support of this notion, we detected expression of the müllerian inhibiting substance in EBs by RT-PCR. While this manuscript was under review, Noce and colleagues reported the successful derivation of male lineage germ cells from ES cells *in vitro*²⁶. Our work corroborates and extends this report by demonstrating the erasure of methylation markers at imprinted loci and the successful fertilization of oocytes by the ES-derived male gametes.

Although EBs may not reflect the precise temporal and spatial features of embryonic development, their ready derivation from ES cells *in vitro* has proven valuable for genetic studies of tissue differentiation, by linking gene deletions or ectopic transgene expression to specific cellular phenotypes. We have demonstrated the *in vitro* differentiation of ES cells into primordial germ cells, which proliferate in response to retinoic acid and give rise to embryonic germ-cell-like clones that undergo erasure of imprints. We are currently using this *in vitro* system to investigate whether EB differentiation sustains a male germ cell niche that enables the proper restoration of male imprints, thereby affording an *in vitro* system to address the genetic and biochemical mechanisms of this fundamental epigenetic modification. As demonstrated by molecu-

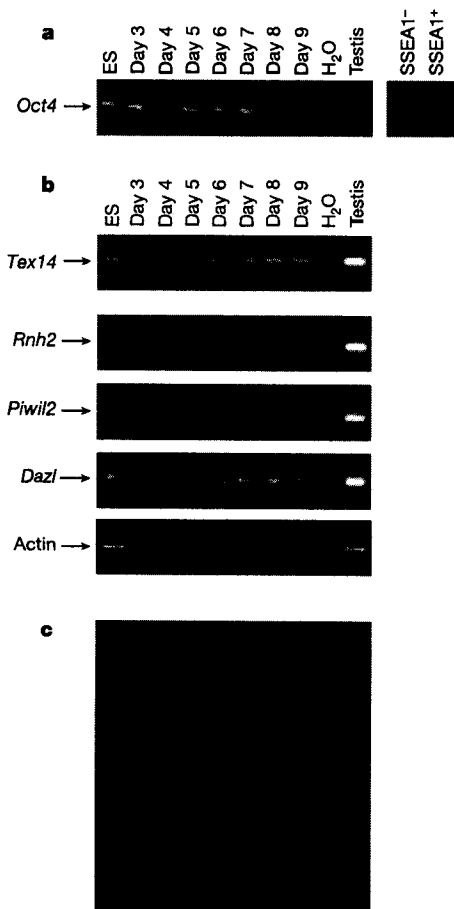


Figure 3 RT-PCR detection of germ-cell-specific genes in SSEA1⁺ cells isolated from developing EBs. **a**, Expression of *Oct4* during EB development (left panel) and in SSEA1⁺ and SSEA1⁻ fractions derived by immunomagnetic sorting. Flow cytometric analysis demonstrates >90% purity in the SSEA1⁺ fraction and >99% purity of the SSEA1⁻ fraction (data not shown). **b**, Expression of *Tex14*, *Rnh2*, *Piwil2* and *Dazl* assayed by RT-PCR. Actin served as a control. **c**, Immunohistochemistry of germ cell and haematopoietic development in day 7 EBs. EBs were cryosectioned and stained with CD41-fluorescein isothiocyanate (green) to mark early haematopoietic cells and SSEA1-PE (red) to visualize germ cell development.

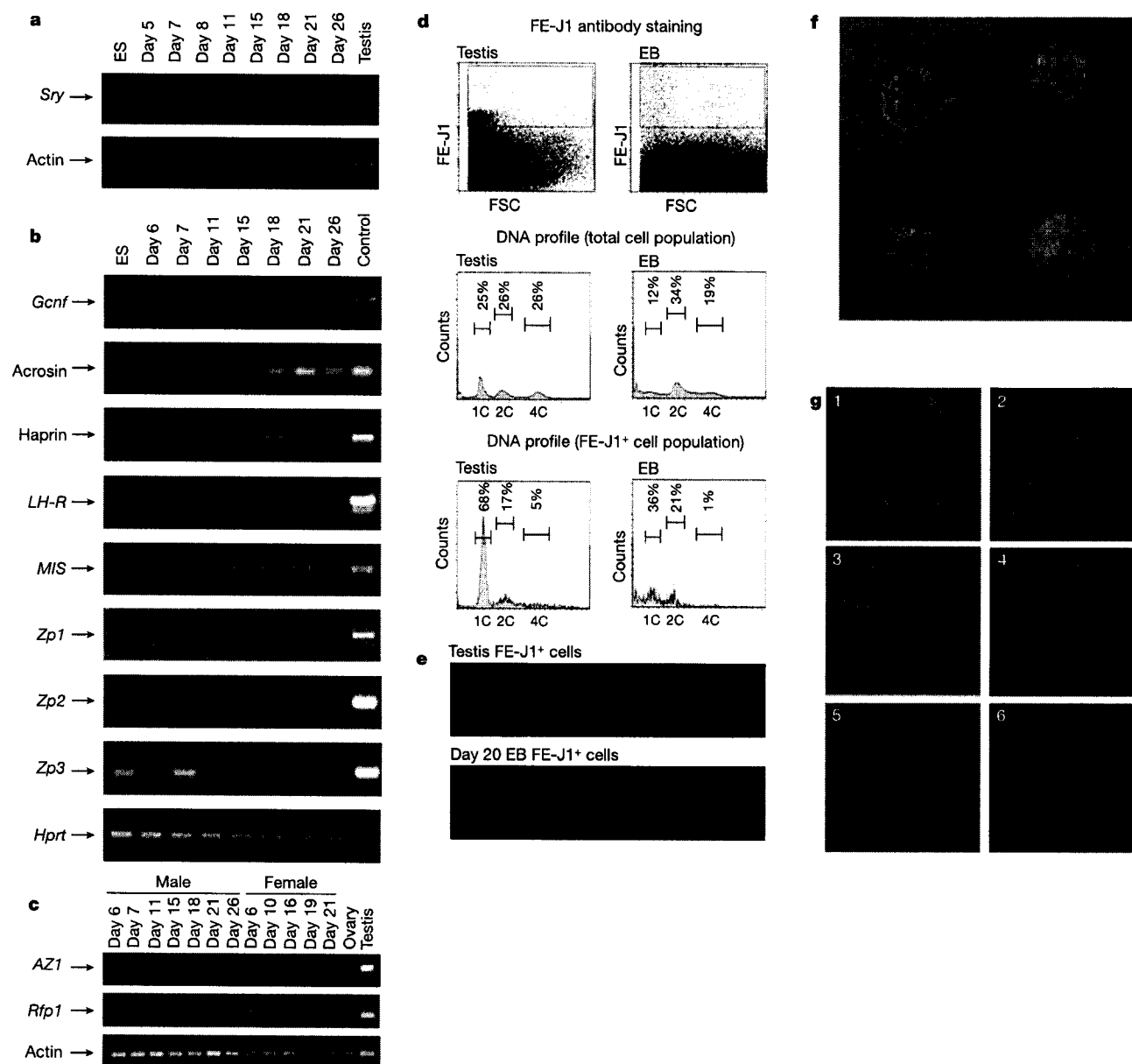


Figure 4 EBs support differentiation of haploid male germ cells that support fertilization of oocytes. **a**, Expression of *Sry* in developing EBs analysed by RT-PCR (top panel). **b**, Expression of genes regulating male germ cell specification during EB development. *Gcnf* helps restrict *Oct4* expression to the germ line¹⁹. *Acrosin*, acrosomal protein with a role in oocyte interaction²⁰; *haprin*, haploid germ-cell-specific RING finger protein; *LH-R* and *MIS*, markers for Leydig and Sertoli cells, respectively; *Zp1–3*, proteins of the oocyte zona pellucida. *Hprt* served as a control. **c**, Expression of male germ-cell-specific genes in male or female EBs assayed by RT-PCR. Control RNA from ovary or testis is shown. **d**, Identification of haploid male germ cells in adult testis (left) or day 13 EBs (right) by immunostaining with the FE-J1 antibody²⁴. Cells were stained with the DNA-binding dye Hoechst 33342 to reveal DNA content. 1C, haploid; 2C, G0/G1 phase; 4C, G2/M phase.

e, Immunohistochemistry of the FE-J1⁺ cells sorted in **d**. FE-J1-PE, red; nuclear Hoechst 33342, blue. **f**, Fluorescence image of blastocysts derived from intracytoplasmic injection of oocytes with EB-derived FE-J1⁺ haploid cells. **g**, FISH analysis of control ES cells and morula stage embryos, with fluorescence signals indicated by arrows. Panels 1 (X and Y chromosome markers) and 2 (autosomal (κ light chain) and Y chromosome markers) show embryos derived from intracytoplasmic oocyte injection of FE-J1⁺ haploid cells. Panels 3 (X and Y chromosome markers) and 4 (autosomal (κ light chain) and Y chromosome markers) show control male ES cells. Panels 5 (X and Y chromosome markers; note two X and absence of Y signals) and 6 (autosomal (κ light chain) and Y chromosome markers; note absence of Y signal) show control female ES cells.

lar analysis, the EBs support a programme of male germ cell differentiation, culminating in the formation of haploid cells that manifest the morphology and fertilization potential of male haploid germ cells of the round spermatid stage. Our report, together with the recent demonstration of oocyte and sperm generation from ES cells^{25,26}, signals a new realm of possibilities for investigating germ cell development, epigenetic reprogramming and germline gene modification.

Methods

Mouse strains and ES cells

C57BL/6-TGN(ACTbEGFP) mice were from Jackson Laboratories²⁷. 129SvEv mice were from Taconic. ES cells were derived from an F₁ cross between C57BL/6-TGN(ACTbEGFP)10sb and 129SvEv.

Cell culture

ES cells were maintained on irradiated MEFs in DME/15% IFS, 0.1 mM non-essential amino acids (GIBCO), 2 mM glutamine, penicillin/streptomycin (GIBCO), 0.1 mM

β -mercaptoethanol, and 1,000 U ml⁻¹ LIF (Invitrogen). For EB differentiation, ES cells were digested with trypsin, collected in EB medium (IMDM/15% FCS, 200 μ g ml⁻¹ iron-saturated transferrin (Sigma), 4.5 mM monothiolglycerol (Sigma), 50 μ g ml⁻¹ ascorbic acid (Sigma) and 2 mM glutamine) and plated for 45 min to allow MEFs to adhere. Non-adherent cells were collected and plated in hanging drops at 200 cells per 30 μ l droplet in an inverted bacterial Petri dish. EBs were collected from the hanging drops at day 3 and transferred into 10 ml EB medium in slowly rotating 10 cm Petri dishes. At day 4, EBs were fed by exchanging half of their spent medium. Cells were collected by collagenase treatment and re-suspension in cell dissociation buffer (Invitrogen).

RT-PCR

RT-PCR amplifications were titrated to be within a linear range of amplification. Primers used are: *Oct4*(f) 5'-GTGGATTCTCGAACCTGGCT-3', *Oct4*(r) 5'-GTCTCCAGACTCCACCTCAC-3'; *stella*(f) 5'-CAGCCGTACCTGTGGAGAACAGAG-3', *stella*(r) 5'-AGCCCTGGGCTCAGAGCTT-3'; *Fgl1*(f) 5'-TTGCTCCGACCATGAACCA-3', *Fgl1*(r) 5'-TGAGCACTTCAGGACCGGA-3'; *Dazl*(f) 5'-GCCAGCACTCAGTCTTCATC-3', *Dazl*(r) 5'-GTTGGAGGCTGCATGTAAGT-3'; *Piwil2*(f) 5'-CCGTCATGAAGAGAGCTCG-3', *Piwil2*(r) 5'-GGAAGCACTCTGTGCTGGAT-3'; *Rnf17*(f) 5'-GACACACAGTCTAACAGAGG-3', *Rnf17*(r) 5'-AGGACAGCAGCATCTACCTT-3'; *Rnh2*(f) 5'-CATAAGTGGCAACGAAGAGC-3', *Rnh2*(r) 5'-GTTACAGGCTGCTACCATCA-3'; *Tdrd1*(f) 5'-GCAGTCTGCTCTGTCAAGG-3', *Tdrd1*(r) 5'-CAGAGCGTGAATCACA TGG-3'; *Tex14*(f) 5'-GAAGCTTGAGCAGGAGGTAG-3', *Tex14*(r) 5'-TTCAAGAGACA CAGACGCCA-3'; *Gcnf*(f) 5'-GTGGAAGACAGGACGACGA-3', *Gcnf*(r) 5'-CCTAC TGGATGATAGTGTGG-3'; *acrosin*(f) 5'-CGGAGTCTACACAGCCACCT-3', *acrosin*(r) 5'-GCATGAGTATGAGGAGGT-3'; *haprin*(f) 5'-CCAGAACATGAGACAGAGAG-3', *haprin*(r) 5'-AGCAACTTCTGAGCATACC-3'; *Hprt*(f) 5'-GCTGGTGAAGAGGACCTCT-3', *Hprt*(r) 5'-CACAGACTAGAACACCTGC-3'; *Sry*(f) 5'-TTACAGCCTGCAG TTGCTC-3', *Sry*(r) 5'-GGTCATAGAAGTGTGTGC-3'; *MIS*(f) 5'-TTGGTGCTAA CCGTGACTT-3', *MIS*(r) 5'-GCAGAGCAGCAACCAAGCGA-3'; *LH-R*(f) 5'-TGCAA CCTCTCAATCTGTC-3', *LH-R*(r) 5'-AGCGTGGCAACCAAGTGGCT-3'; *Zp1*(f) 5'-GAGTGACTGTGTGCCATAG-3', *Zp1*(r) 5'-GCCACACTGGTCTCACTACG-3'; *Zp2*(f) 5'-GCTACACATGACTCTCAC-3', *Zp2*(r) 5'-GGTGACTCACAGTGGCA CTC-3'; *Zp3*(f) 5'-TTGAGCAGAGAGCTCCAGC-3', *Zp3*(r) 5'-CGGTGTGCCTTGT GGATGGTC-3'; *actin*(f) 5'-ACCAACTGGGACGATATGGAGAAGA-3', *actin*(r) 5'-CTCTTTGATGTACGACGATTTC-3'.

Immunomagnetic isolation of SSEA1⁺ cells

Cells collected from EBs were incubated (30 min) with a monoclonal antibody against SSEA1 (Hybridoma bank) at 4 °C in PBS/0.5% BSA. Cells were washed twice with ice-cold PBS/0.5% BSA before addition of immunomagnetic rat anti-mouse IgM beads (Dyna) and were incubated for 1 h at 4 °C with slow rotation. Magnetic separation of SSEA1⁺ beads associated with the cells was performed according to the manufacturer's protocol.

Southern analysis

Genomic DNA was prepared from individual clones of the parental ES cell line or EB-derived embryonic germ cells. For the embryonic germ cell clones, SSEA1⁺ cells were isolated at different days of EB development and cells were grown in the presence of retinoic acid for 7 days followed by 2 days of culture without retinoic acid. Individual clones were isolated and expanded on gelatinized tissue culture plastic in the presence of LIF to remove feeder cells. Genomic DNA was isolated and digested with *PvuII* and *MluI* for the detection of *Igf2r* methylation, or with *SacI* and *HhaI* for the analysis of *H19* imprints. DNA was separated on a 0.7% agarose gel and Southern blots were generated by standard methods. Filters were hybridized with a probe covering region 2 of the *Igf2r* receptor (pPP4).

FACS analysis

Testicular and EB-derived cell suspensions were obtained by enzymatic digest of the tissue. Briefly, cells were incubated at 37 °C for 15 min with digest buffer (0.1% collagenase IV, 0.2% hyaluronidase and 50 U ml⁻¹ DNase (all Sigma)). Cells were then dissociated using cell dissociation buffer (Invitrogen), collected by centrifugation and a second digest was performed. Cell clumps were removed using a 70 μ m strainer and cells were re-suspended in ice-cold RPMI plus 0.5% FBS. Cells were incubated with FE-J1, a haploid male germ-cell-specific antibody (Hybridoma bank²⁴), for 30 min at 4 °C. The cells were washed twice with RPMI/0.5% FBS and incubated with phycoerythrin (PE)-conjugated rat anti-mouse IgM for 30 min at 4 °C. Cells were washed twice with RPMI/0.5% FBS, re-suspended in RPMI/0.5% FBS containing 2.5 μ g ml⁻¹ cytochalasin B (Sigma) and sorted on a Becton-Dickinson FACSCalibur.

Oocyte injections

Eight-ten-week-old B6D2F1/J mice (Jackson Laboratories) were used as oocyte donors. EB-derived donor cells were re-suspended in KSOM (Specialty Medium) containing 10% (w/v) polyvinyl alcohol (Sigma) and 0.01% (w/v) bovine serum albumin (Sigma). Intracytoplasmic injection into cumulus-free oocytes was carried out in H-KSOM containing 5 μ g ml⁻¹ cytochalasin B (Sigma) and 3% (w/v) sucrose at room temperature using a Nikon Eclipse TE300 microscope equipped with Narashige hydraulic micromanipulators and Hoffman modulation contrast. The injected oocytes were washed five times in KSOM to remove the cytochalasin. Reconstructed embryos were activated either in KSOM containing 10 μ M calcium ionophore A23187 (Sigma) for 5 min, followed by 2 mM 6-dimethylaminopurine (Sigma) in KSOM at 37 °C in 5% CO₂ for 3 h, or for 5 h in Ca²⁺-free medium containing 10 mM SrCl₂. Embryos were then washed five times in KSOM. Embryos were cultured in KSOM at 37 °C in 5% CO₂.

DNA-FISH

DNA-FISH was performed as described²⁸ with minor modifications. The zona was removed from embryos using acid tyrodase, and the morulae were incubated in a small drop of 0.075 M KCl on a slide for 5 min. Embryos were fixed by replacing the KCl with 3:1 methanol:acetic acid. Cells were permeabilized in 0.5% triton/PBS for 10 min, washed twice in PBS and dehydrated in 70%, 90% and 100% ethanol. Bacterial artificial chromosome (BAC) clones used for FISH analysis were from BACPAC resources and from Invitrogen. Autosomal sequences were detected with BAC RP23-20P21, which is specific for the κ light chain locus of immunoglobulin. X chromosomal sequences were detected with BAC CT7-22804, which is specific for the *Irak1* locus. Y chromosomal sequences were detected with BAC RP24-507D23, which is specific for Y chromosomal repeats.

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T-cell priming by dendritic cells in lymph nodes occurs in three distinct phases

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Primary T-cell responses in lymph nodes (LNs) require contact-dependent information exchange between T cells and dendritic cells (DCs). Because lymphocytes continually enter and leave normal LNs, the resident lymphocyte pool is composed of non-synchronized cells with different dwell times that display heterogeneous behaviour in mouse LNs *in vitro*^{1–3}. Here we employ two-photon microscopy *in vivo* to study antigen-presenting DCs and naive T cells whose dwell time in LNs was synchronized. During the first 8 h after entering from the blood, T cells underwent multiple short encounters with DCs, progressively decreased their motility, and upregulated activation markers. During the subsequent 12 h T cells formed long-lasting stable conjugates with DCs and began to secrete interleukin-2 and interferon- γ . On the second day, coinciding with the onset of proliferation, T cells resumed their rapid migration and short DC contacts. Thus, T-cell priming by DCs occurs in three successive stages: transient serial encounters during the first activation phase are followed by a second phase of stable contacts culminating in cytokine production, which makes a transition into a third phase of high motility and rapid proliferation.

Naive T cells recirculate continually between the blood and LNs to search for antigen⁴. The intranodal encounter of a peptide-major histocompatibility complex (MHC) complex that is recognized by a T-cell antigen receptor (TCR) will only result in full-fledged T cell activation upon co-stimulation provided by mature DCs. These professional antigen-presenting cells collect antigens in peripheral tissues and migrate to LNs through lymph vessels. T-cell priming by DCs induces activation markers, cytokine secretion, and proliferation. Several reports have analysed the dynamics of T-cell–DC interactions in excised LNs, but the methods and results have been variable and it is unknown how the absence of lymph and blood flow or innervation influences T-cell–DC interactions^{1–3,5}. There is therefore still no comprehensive description of what happens in a truly physiological setting when naive T cells enter LNs that contain antigen-presenting DCs.

We have used two-photon microscopy⁶ to study lymphocyte migration and interactions with DCs within popliteal LNs of anaesthetized mice. Our preparation preserved physiological blood and lymph flow⁷, whereas, in our hands, lymph flow was

compromised when we attempted to adapt the well-established inguinal lymph-node preparation to two-photon imaging^{8,9} (not shown). Recipient mice received footpad injections of fluorescent DCs, which entered lymph vessels and accumulated in the popliteal LN during the following day. Differentially tagged TCR transgenic CD8⁺T cells were injected intravenously 18 h later (Fig. 1a). T cells homed rapidly through high endothelial venules (HEVs) into popliteal LNs⁷, where they constituted 1–2% of all CD8⁺ cells 2 h after injection. At this point, further lymphocyte homing was blocked by the injection of anti-L-selectin¹⁰. This ensured that all imaged T cells had entered popliteal LNs during the initial 2-h window and enabled us to study synchronized resident cells during the subsequent 2 days.

Initially, we examined the phenotype of injected DCs that migrated to draining LNs (Fig. 1b). Immature splenic CD11c⁺ DCs from CD45.1⁺ donors were injected into CD45.2⁺ congenic recipients. Because immature DCs express little or no CCR7—a chemokine receptor required for DC migration into lymphatics and within LNs¹¹—we co-injected lipopolysaccharide (LPS; 10 ng), which induces DC maturation and CCR7 expression¹². Consequently, DCs recovered from popliteal LNs were mature and either CD11b⁺CD8 α [–] or CD11b[–]CD8 α ^{low/–}. CD8 α ^{high} input DCs were rarely detected in LNs. This might reflect an inability of CD8 α ^{high} DCs to migrate to LNs, or the downregulation of CD8 α during transit.

To examine DCs *in situ*, we administered footpad injections of red fluorescent DCs and analysed their behaviour within popliteal LNs. After 20 h a fraction of migrated DCs (about 20–50%) resided in the subcapsular sinus and the superficial cortex. The remainder congregated in distinct regions within the deep cortex, where they localized with homed T cells (Supplementary Fig. S1a). A reason for the non-uniform distribution of DCs in the T cell area became apparent after fluorescein isothiocyanate–dextran injection to delineate blood vessels. In line with recent observations¹³, many DCs lined up around HEVs in strategic positions to interact with newly homed T cells (Supplementary Fig. S1b; Supplementary Information 2, 3). This distribution pattern became less apparent at day 2 or 3 after injection (Supplementary Information 4).

One day after injection, DCs in T-cell areas were remarkably motile, whether they presented antigen to T cells or not (Fig. 1c–g; Supplementary Information 5). The median three-dimensional (3D) instantaneous velocity was 6.6 $\mu\text{m min}^{-1}$ (Fig. 1e), which is in good agreement with measurements of two-dimensional velocities in explanted LNs³. Individual DCs followed random paths without apparent directional bias. Even sessile DCs constantly extended and retracted dendrites and pseudopods. The motility of both antigen-pulsed and control DCs was highest between 2 and 8 h after T-cell transfer (20–26 h after DC injection) and decreased over time (Fig. 1f, g; Supplementary Information 5). This progressive decrease in motility might reflect DC ‘exhaustion’, which has been proposed to occur after LPS activation¹⁴.

Migratory dynamics in LNs were closely dependent on physiological conditions: T cells and DCs stopped migrating and assumed a round shape within minutes after cardiovascular arrest (Supplementary Information 6), even with a tissue temperature maintained at 36 °C, a requirement for interstitial T-cell migration². In living animals without DC injection, intranodal T cells moved rapidly, reaching peak 3D velocities of 40 $\mu\text{m min}^{-1}$ (Fig. 2a–c; Supplementary Information 7). In LNs draining DC-injected footpads, 3D velocities were equivalent (Fig. 2c, d) but T cells turned at steeper angles and covered a smaller volume of the paracortex, resulting in decreased motility coefficients, even when DCs carried no antigen (Fig. 2e, f). Mean T-cell displacement plots in the antigen-containing LNs revealed a plateau at about 15 μm displacement (Fig. 2e), indicating confined motility¹⁵. Thus, DC plus LPS injections induced LN paracortex partitioning into smaller compartments in which incoming T cells were retained, perhaps owing

Differentiation of Embryonic Stem Cells After Transplantation Into Peritoneal Cavity of Irradiated Mice and Expression of Specific Germ Cell Genes in Pluripotent Cells

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ABSTRACT

Permanent embryonic stem cell lines (ES cells) are considered as one of the most promising cellular sources for regenerative medicine. ES cells have a high proliferative potency and ability to differentiate into all kinds of somatic and germ cells. However, transplantation of undifferentiated ES cells into adult recipient tissue results in the formation of teratomas. To understand the mechanisms underlying self-renewal and determination of pluripotent cells, we investigated differentiation potencies of undifferentiated ES cells and differentiating embryoid bodies (EB). ES cells and EBs growing on acetate-cellulose membranes were transplanted into the peritoneal cavity of irradiated mice. Behavior and differentiation of transplanted cells were studied within 1, 2, 3, and 6 weeks after transplantation. No differences in the cell composition were found in the teratomas formed by ES cells and differentiating EBs. The pattern of expression of the genes specific for pluripotent and germ cells was studied in all types of experimental teratomas. The expression of *oct4*, *stella*, *fragilis* was detected in the teratomas, but *nanog* was not expressed. We conclude that pluripotent cells are retained in the experimental teratomas formed after transplantation of ES cells and EBs but the pattern of expression of the studied genes underwent changes.

PLURIPOTENT embryonic stem (ES) cell lines can be expanded for prolonged period of culture, maintain a normal karyotype, and have potency to generate all cell types. Therefore they represent a unique source for the repair of diseased or damaged tissues in human body.

Predominant ES cell differentiation into required cell types is only possible under special conditions of co-culture and 3D-culture^{1,2} and in the presence of exogenous growth and differentiation factors.³⁻⁷ Analysis of gene expression pattern during differentiation of ES cells into several somatic cell types demonstrates certain similarities with the processes of histogenesis.^{3,6-8} Several authors reported successful transplantation of the precursors of neurons, hepatocytes, and endothelial cells derived from ES cells and isolated using a fluorescent-activated cell sorting (FACS). Their behavior, duration of preservation in recipient tissues, and activity of certain marker genes have been studied.^{6,9,10} On the other hand, when pluripotent stem cells are introduced into adult histocompatible or immunocompromised mice, they form teratomas or teratocarcinomas that include derivatives of all germ layers.^{11,12} Analysis of interactions between pluripotent cells and differentiated

cells of the recipient animals, identification of the factors that drive differentiation of ES cells in vivo can help in understanding the mechanisms involved in the determination of cell fate during normal development and oncogenesis. These data are important for development of effective and safe stem cells technologies for prospective clinical treatment.

It was shown that two genes, *oct4* and *nanog*, were involved in the regulation of pluripotent state of ES cells, inner mass cells of blastocyst, early epiblast, and developing germ cells.¹³⁻¹⁵ These genes are also expressed in other pluripotent cell lines: embryonic germ cells and embryonal carcinoma stem cells.¹⁶ Expression of *stella* and *fragilis* genes was identified in founders primordial

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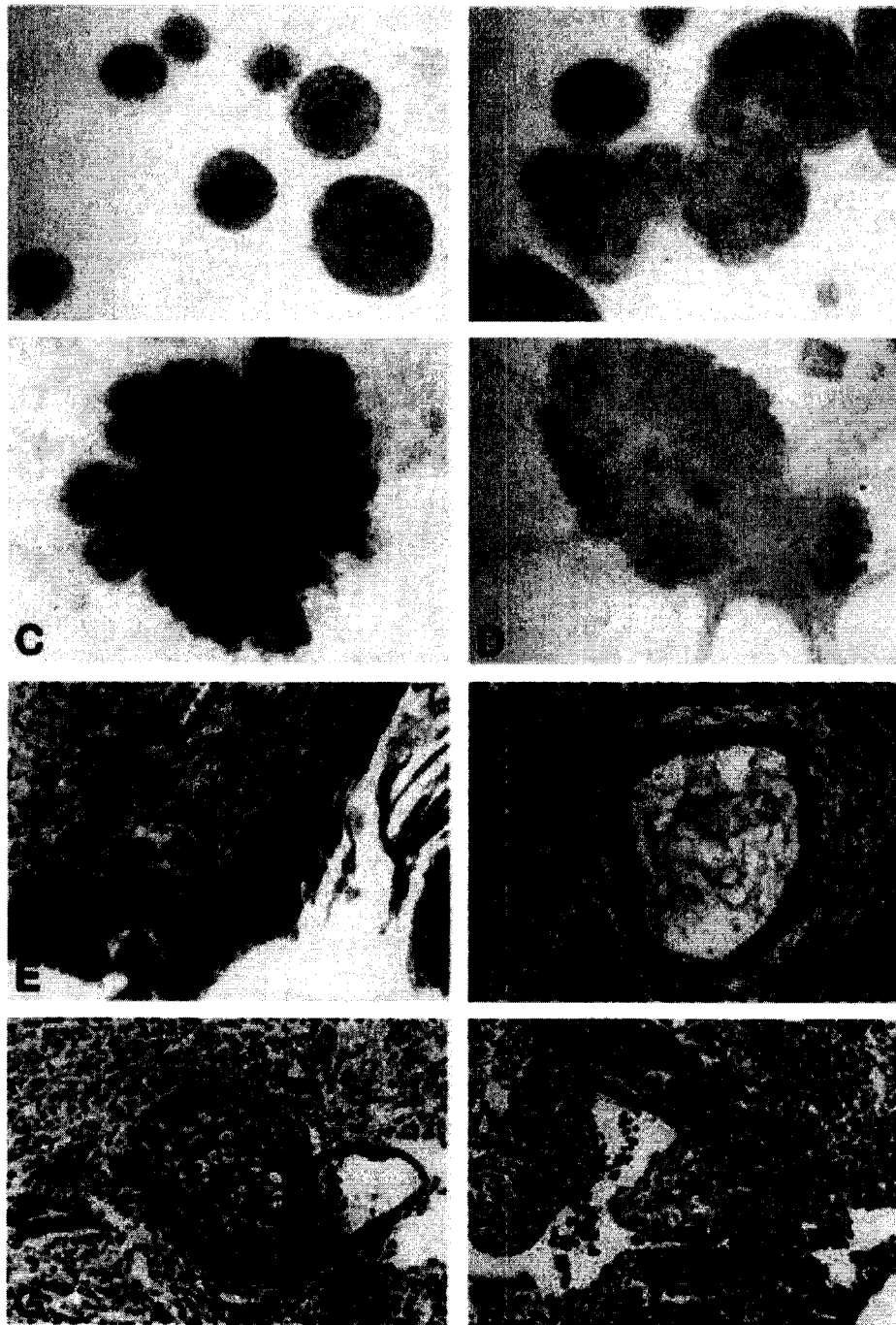


Fig 1. Differentiation of pluripotent populations of ES cell colonies and differentiating EBs *in vivo*. The activity of alkaline phosphatase in ES cells (**A,C**) and EBs (**B,D**) prior and within 1 week after transplantation, respectively. Sections through teratomas contain a variety of tissue types derived of the three germ layers: keratinizing epithelia (**E**), secretory epithelial cyst (**F**), and cartilage (**G**). Lacunar blood vessel in teratoma (**H**). Original magnification: $\times 100$.

germ cells, as well as in ES cells.¹⁷ However, the data on the fate of pluripotent ES cells and on the mechanisms of their destination after transplantation into adult animals are very limited. It is not known how pluripotent cells that were retained in teratomas change their phenotype, potencies, and expression pattern.

We present herein the data about the differentiation potencies of ES cells and differentiating EBs and the expression of specific pluripotent and germ cells genes in teratomas formed by these pluripotent populations after transplantation into peritoneal cavity of irradiated mice.

MATERIALS AND METHODS

For the experiments, we used the murine embryonic stem cell line R1 kindly provided by Dr A. Nagy (Mount Sinai Hospital, Toronto, Canada). Undifferentiated ES cells growing on a feeder layer of the mouse primary embryonic fibroblasts inactivated by mitomycin C (10 μ g/mL; Sigma, St. Louis, Mo, USA) as well as the differentiating EBs (3 to 5 days after plating) were cultured on gelatinized acetate cellulose (AC) membranes (Millipore), as described previously.¹⁸ Three days after the attachment to the substrate, the membranes with ES cell colonies or EBs were implanted into the peritoneal cavity of hybrid mice (CBA \times C57B1) F1 irradiated 3 to 5 days before at 5.72 Gy on a RUM-17 device (210 kV, 15 mA, filters: 0.55 mm Cu + 1.0 mm Al, 0.78 Gy/min dose power, 47 cm distance from the source). The AC membranes with the primary embryonic fibroblasts inactivated by mitomycin C or empty gelatin-coated membranes were implanted in the control animals.

The membranes with cells and fragments of tissues were isolated after 1, 2, 3, and 6 weeks after the implantation for histologic study of the total preparations and paraffin sections. The material was fixed by 4% paraformaldehyde (Sigma) for 30 minutes. The preparations were stained by azur-eosin or hematoxylin and eosin. The activity of alkaline phosphatase was detected in fixed ES cells and EBs after incubation in a solution of naphthol AS-BI-phosphate and Fast Red dye Texas Red (Sigma) at 37°C for 1 hour.

For analysis of gene expression, total RNA was isolated using the mixture RNazol B (Ginna-Biotecs, Friendswood). A fraction of mRNA was obtained from total RNA using a kit for mRNA isolation with magnetic carriers covered by oligo-(dT)-cellulose (Dynabeads mRNA purified kit, Dynal, Norway). cDNA copies were synthesized on mRNA templates from ES cells and differentiating EBs with the help of revertase SuperScript using oligo-(dT)₁₈ and random primers according to the producer's protocol (Gibco). The synthesized cDNA libraries were normalized by evaluation of hprt gene expression level. The levels of expression of the regulatory genes in ES cells, EBs, and experimental teratomas were quantitatively estimated by PCR analysis using specific primers constructed on the basis of the data on the structure of the genes we studied. The primers used were oct4 (f) 5'-TGGAGACTTTGC-AGCCTGAG-3', oct4 (r) 5'-CATACTCTTCTCGTTGGGAAT-A-3'; nanog (f) 5'-ATGAGTGTGGGTCTTCTCGGT-3', nanog (r) 5'-TATTCACCTGGAGTCACA-3'; stella (f) 5'-CAGCCGTACCTGTGGAGAAC-3', stella (r) 5'-AGCCCTGGGGCC-TCACAGCTT-3'; fragilis (f) 5'-TGCTCCGCACCATGAACC-AC-3', fragilis (r) 5'-GTGAAGCACTTCAGGACCGG-3'.

RESULTS AND DISCUSSION

Comparative morphologic analysis of the differentiation of pluripotent cells in the ES cell colonies and EBs in the peritoneal cavity of irradiated mice revealed cell clusters on the implanted membranes resulting from proliferation and migration of the transplanted cells 1 and 2 weeks later in each experimental series. By this time, the colonies of undifferentiated ES cells merged and assumed asymmetric shape, while outgrowths of the EBs were enlarged. No morphologically differentiated cell types were revealed in the cell clusters, and the activity of alkaline phosphatase was detected in most cells (Fig 1A-D). In addition to ES cells, numerous fibroblast-like cells of the recipient encapsulating the foreign body were visualized on the mem-

branes. These fibroblast-like cells were also presented on the AC membranes isolated from the abdominal cavity of control animals. After 3 weeks of the experiment, the teratomas developed in all experimental animals in the region of contact between the membrane and small intestinal loops; however, no tumors were found in other tissues. After 6 weeks, the teratomas formed by ES cells were considerably larger than after 3 weeks.

Histologic analysis revealed various types of differentiated cells in the experimental teratomas, including ectodermal (keratinizing epithelium and neural ganglion cells), mesodermal (hyaline cartilage and adipocytes), and endodermal (epithelium of intestinal and respiratory types) lineages (Fig 1D-E). Lacunar blood vessels (most likely of the recipient origin) and vast hemorrhagic zones were seen in the teratomas (Fig 1H). Apparently, the observed necrotic changes in the teratomas developed as a result of immune conflict with the immune system of the recipient. Our experiments have revealed no considerable changes in the types of differentiations of pluripotent populations of the ES cell colonies and the EBs, although the differentiating EBs attached to the substrate represented heterogeneous populations including pluripotent, differentiated, and committed cells.

Expression of pluripotent and germ cell-specific genes oct4, nanog, stella, and fragilis was evaluated using normalized cDNA libraries constructed by RT-PCR from undifferentiated ES cells, differentiating embryoid bodies, all types of experimental teratomas, and primary embryonic fibroblasts (feeder cells). The pattern of expression of the studied genes was identical for ES cells and differentiating embryoid bodies prior to transplantation. All four genes were expressed in the pluripotent cells. In contrast, we identified expression of the oct4, stella, and fragilis; nanog expression was not detected in all types of experimental teratomas. The levels of expression of the detected genes were approximately equal. Our findings suggest that the

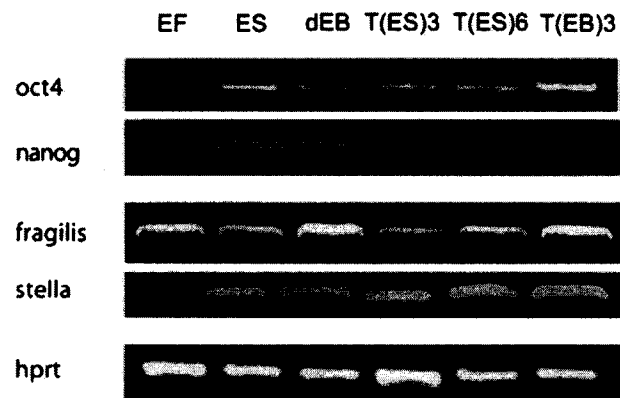


Fig 2. The pattern of expression of germ cell-specific genes in ES cells (ES), differentiating EB(dEB) in the experimental teratomas formed by ES cells [T(ES)] and EBs [T(EB)] within 3 and 6 weeks after transplantation. EF, feeder cells. cDNA libraries were diluted 1:5 for analysis of oct4 expression.

populations of ES cells and differentiating EBs have the similar developmental potencies for differentiation into derivatives of three germ layers and that pluripotent cells retained in the experimental teratomas have the same expression pattern of the known pluripotent and germ cell-specific genes (Fig 2). Note that similar pattern of expression of the studied genes were described in migrating primordial germ cells in the mouse embryo.¹⁷

Hence, pluripotent cells are retained in the teratomas formed after transplantation of ES cells and EBs but the expression pattern changed in them as compared to initial pluripotent populations.

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